=> d his

(FILE 'HOME' ENTERED AT 10:16:01 ON 16 MAY 2003)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, SCISEARCH, CANCERLIT, TOXCENTER' ENTERED AT 10:16:23 ON 16 MAY 2003

L1 1743 S HEPARANASE

L2 0 S L1 AND PARALOG

L3 182 S L1 AND HUMAN HEPARANASE

L4 2 S L3 AND (SPLICE VARIANT OR PARALOG)

L5 2 DUP REM L4 (0 DUPLICATES REMOVED)

=>

=> dup rem 14 PROCESSING COMPLETED FOR L4 2 DUP REM L4 (0 DUPLICATES REMOVED)

=> d 15 ibib ab 1-2

ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2002:51645 CAPLUS

DOCUMENT NUMBER:

136:97379

TITLE:

Protein and cDNA sequences of a second human

heparanase, and splice

variants thereof, with a predominant

expression in skeletal muscle, heart and pancreas

INVENTOR(S):

David, Guido; Duerr, Joachim

PATENT ASSIGNEE(S):

Vlaams Interuniversitair Instituut voor Biotechnologie

Vzw, Belg.

SOURCE:

PCT Int. Appl., 54 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA	PATENT NO.					DATE					CATIO			DATE			
	2002	0046	45	A2	2									2001	0712		
	₩:	AE, CO, GM, LS, RO, UZ, GH, DE,	AG, CR, HR, LT, RU, VN, GM, DK,	AL, CU, HU, LU, SD, YU, KE, ES,	AM, CZ, ID, LV, SE, ZA, LS,	AT, DE, IL, MA, SG, ZW, MW, FR,	AU, DK, IN, MD, SI, AM, MZ, GB,	DM, IS, MG, SK, AZ, SD, GR,	DZ, JP, MK, SL, BY, SL, IE,	EC, KE, MN, TJ, KG, SZ, IT,	EE, KG, MW, TM, KZ, TZ, LU,	ES, KP, MX, TR, MD, UG, MC,	FI, KR, MZ, TT, RU, ZW, NL,	GB, KZ, NO, TZ, TJ, AT, PT,	GD, LC, NZ, UA, TM BE, SE,	GE, LK, PL, UG,	GH, LR, PT, US,
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG PRIORITY APPLN. INFO.: AB The present invention provides a novel human protein and its splicing variants, which has heparanase activity. The present invention relates to the field of carbohydrates and more specifically to the field of heparan sulfate proteoglycans. Several splice variants of said gene have been identified with a specific expression pattern in skeletal muscle, heart and pancreas.										ng							

ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS L5

ACCESSION NUMBER:

2001:472900 CAPLUS

DOCUMENT NUMBER:

135:73335

TITLE:

A human heparanase sequence homolog and splice variants and

their possible therapeutic use in the control of

invasive cell proliferation

INVENTOR (S):

Mckenzie, Edward Alexander; Stamps, Alasdair Craig; Terrett, Jonathan Alexander; Tyson, Kerry Louise

Oxford Glycosciences (Uk) Ltd., UK

PATENT ASSIGNEE(S): SOURCE:

PCT Int. Appl., 97 pp.

CODEN: PIXXD2

_ _ _ _ _ _

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.

KIND DATE

APPLICATION NO. DATE

WO 2000-GB4963 20001221 WO 2001046392 20010628 A2 WO 2001046392 Α3 20011206 AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG EP 1240313 .240313 A2 20020918 EP 2000-985677 20001221 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR A1 20030501 20020621 US 2002-177245 US 2003083254 A 19991222 PRIORITY APPLN. INFO.: GB 1999-30392 GB 2000-8713 20000407 W 20001221 WO 2000-GB4963

AB A human sequence homolog of **heparanase** and a no. of variants that can arise from alternative splicing are described. The protein may play a role in the control of heparan-dependent invasive cell growth in a no. of pathologies and may therefore be a target for therapeutics. Identification of an EST for a **heparanase** homolog in a com. sequence database, PCR cloning of a cDNA and anal. of tissue distribution of the mRNA are reported.

(FILE 'HOME' ENTERED AT 09:39:05 ON 16 MAY 2003)

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DDFB, DDFU, DGENE, DRUGB, DRUGLAUNCH, DRUGMONOG2, ...' ENTERED AT 09:39:21 ON 16 MAY 2003

SEA (HUMAN HEPARANASE II OR HUMAN HEPARANASE-2 OR HNHP1)

- 1 FILE BIOTECHABS
- 1 FILE BIOTECHDS
- 3 FILE CAPLUS
- 63 FILE DGENE
- 1 FILE IFIPAT
- 1 FILE USPATFULL
- L1 QUE (HUMAN HEPARANASE II OR HUMAN HEPARANASE-2 OR HNHP1)

FILE 'DGENE, CAPLUS, BIOTECHDS, IFIPAT, USPATFULL, ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHNO' ENTERED AT 09:43:21 ON 16 MAY 2003

L2 69 S L1

L4

L6

L3 68 DUP REM L2 (1 DUPLICATE REMOVED)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, SCISEARCH, CANCERLIT' ENTERED AT 09:45:52 ON 16 MAY 2003

1601 S HEPARANASE

L5 174 S (HUMAN HEPARANASE)

- 4 S (HEPARANASE-2 OR HEPASRANASE II)
- L7 2 S L5 AND (HEPARANASE-2 OR HEPASRANASE II)
- L8 10 S L5 AND (VARIANT OR MUTANT OR SPLICE VARIANT)
- L9 5 DUP REM L8 (5 DUPLICATES REMOVED)

=> d 19 ibib ab 1-5

L9 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER:

2002:51645 CAPLUS

DOCUMENT NUMBER:

136:97379

TITLE:

Protein and cDNA sequences of a second human

heparanase, and splice

variants thereof, with a predominant

expression in skeletal muscle, heart and pancreas

David, Guido; Duerr, Joachim

PATENT ASSIGNEE(S):

Vlaams Interuniversitair Instituut voor Biotechnologie

Vzw, Belg.

SOURCE:

PCT Int. Appl., 54 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

INVENTOR(S):

Patent English

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

P	PATENT NO.							APPLICATION NO. DATE										
	10	2002004645		A:	2	2002	0117	WO 2001-EP8094 20010712										
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		W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	AΞ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,
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L9 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER:

2001:472900 CAPLUS

DOCUMENT NUMBER:

135:73335

TITLE:

A human heparanase sequence homolog and splice variants and

their possible therapeutic use in the control of

invasive cell proliferation

INVENTOR(S):

Mckenzie, Edward Alexander; Stamps, Alasdair Craig; Terrett, Jonathan Alexander; Tyson, Kerry Louise

PATENT ASSIGNEE(S):

Oxford Glycosciences (Uk) Ltd., UK PCT Int. Appl., 97 pp.

SOURCE: PCT Int. Appl CODEN: PIXXD2

Patent

DOCUMENT TYPE:

English

LANGUAGE: FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLI	CATION NO.	DATE	
WO 2001046392	A2	20010628	WO 20	00-GB4963	20001221	
WO 2001046392	A3	20011206				
W: AE, A	G, AL, AM	, AT, AU,	AZ, BA, BB,	BG, BR, BY	, BZ, CA,	CH, CN,
CR, CI	J, CZ, DE	, DK, DM,	DZ, EE, ES,	FI, GB, GD	, GE, GH,	GM, HR,

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HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
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               SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,
                YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
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                                              EP 2000-985677 20001221
                           A2 20020918
      EP 1240313
           R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
                                                                          20020621
                          A1 20030501
                                                     US 2002-177245
      US 2003083254
                                                 GB 1999-30392 A 19991222
PRIORITY APPLN. INFO.:
                                                                      A 20000407
                                                 GB 2000-8713
                                                 WO 2000-GB4963 W 20001221
```

A human sequence homolog of heparanase and a no. of variants AB that can arise from alternative splicing are described. The protein may play a role in the control of heparan-dependent invasive cell growth in a no. of pathologies and may therefore be a target for therapeutics. Identification of an EST for a heparanase homolog in a com. sequence database, PCR cloning of a cDNA and anal. of tissue distribution of the mRNA are reported.

ANSWER 3 OF 5 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER:

2001:12473 CAPLUS

DOCUMENT NUMBER:

134:96257

TITLE:

Protein and cDNA sequences of a novel human

heparanase gene hnhp1 and its splicing

variants

INVENTOR(S):

PATENT ASSIGNEE(S):

SOURCE:

Pecker, Iris; Michal, Israel; Itzhaki, Hanan Insight Strategy & Marketing Ltd., Israel

PCT Int. Appl., 67 pp. CODEN: PIXXD2

DOCUMENT TYPE:

LANGUAGE:

Patent English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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PATENT NO. KIND DATE
                                                           APPLICATION NO. DATE
       WO 2001000643 A2 20010104 WO 2000-IL358 20000619
              W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
                     CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
              CR, CU, CZ, DE, DK, DM, DZ, EE, ES, F1, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                    A1 20020612
                                                                  EP 2000-937164 20000619
       EP 1212341
                  AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL
                                                                    JP 2001-507050
        JP 2003503070
                               T2 20030128
                                                                                                20000619
                                                                   NO 2001-5526 20011112
        NO 2001005526 A 20011218
                                                                US 1999-140801P P 19990625
PRIORITY APPLN. INFO.:
                                                                WO 2000-IL358 W 20000619
```

The invention provides protein and cDNA sequences of a novel human AB heparanase gene hnhp1 and two variants resulted from alternative splicing. The longest clone is 2060 nucleotide long and it contains an open reading frame of 1776 nucleotides, which encodes a polypeptide of 592 amino acids, with a calcd. mol. wt. of 66.5 kDa. two shorter forms contain an in frame deletion as a result of alternative splicing, one is 162 nucleotides (nt473-634) corresponding to amino acids 150-203, and one is 336 nucleotides (nt473-808) corresponding to amino

acids 150-261. The hnhp1 gene is mapped to chromosome 10, next to the marker SHGC-57721. The tissue distribution of hnhp1 transcripts is detd. The invention also relates to constructing hnhpl gene expression vector to produce recombinant proteins in mammalian cells, which may have heparanase or other glycosyl hydrolase activity, its antibodies, and antisense oligonucleotide and ribozymes for gene modulation and therapeutic use.

THERE ARE 52 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 52

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 4 OF 5 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 1 1.9

ACCESSION NUMBER: DOCUMENT NUMBER:

2001:50122 BIOSIS PREV200100050122

TITLE:

Identification of active-site residues of the pro-metastatic endoglycosidase heparanase.

AUTHOR (S):

Hulett, Mark D.; Hornby, June R.; Ohms, Stephen J.; Zuegg,

Johannes; Freeman, Craig; Gready, Jill E.; Parish,

Christopher R. (1)

CORPORATE SOURCE:

(1) Division of Immunology and Cell Biology, John Curtin School of Medical Research, Australian National University,

Canberra, ACT, 2601: Christopher.Parish@anu.edu.au

Australia

SOURCE:

Biochemistry, (December 26, 2000) Vol. 39, No. 51, pp.

15659-15667. print. ISSN: 0006-2960.

DOCUMENT TYPE:

General Review

LANGUAGE:

English

English SUMMARY LANGUAGE:

Heparanase is a beta-D-endoglucuronidase that cleaves heparan sulfate (HS) and has been implicated in many important physiological and pathological processes, including tumor cell metastasis, angiogenesis, and leukocyte migration. We report herein the identification of active-site residues of human heparanase. Using PSI-BLAST and PHI-BLAST searches of sequence databases, similarities were identified between heparanase and members of several of the glycosyl hydrolase families (10, 39, and 51) from glycosyl hydrolase clan A (GH-A), including strong local identities to regions containing the critical active-site catalytic proton donor and nucleophile residues that are conserved in this clan of enzymes. Furthermore, secondary structure predictions suggested that heparanase is likely to contain an (alpha/beta) 8 TIM-barrel fold, which is common to the GH-A families. On the basis of sequence alignments with a number of glycosyl hydrolases from GH-A, Glu225 and Glu343 of human heparanase were identified as the likely proton donor and nucleophile residues, respectively. The substitution of these residues with alanine and the subsequent expression of the mutant heparanases in COS-7 cells demonstrated that the HS-degrading capacity of both was abolished. In contrast, the alanine substitution of two other glutamic acid residues (Glu378 and Glu396), both predicted to be outside the active site, did not affect heparanase activity. These data suggest that heparanase is a member of the clan A glycosyl hydrolases and has a common catalytic mechanism that involves two conserved acidic residues, a putative proton donor at Glu225 and a nucleophile at Glu343.

ANSWER 5 OF 5 CANCERLIT

93696451 CANCERLIT ACCESSION NUMBER:

DOCUMENT NUMBER:

93696451

TITLE:

The molecular cloning and characterization of human heparanase cDNA and the immunochemical localization

of heparanase in metastatic melanomas.

AUTHOR:

CORPORATE SOURCE:

Univ. of Texas H.S.C. at Houston Grad. Sch. of Biomed. Sci.

Diss Abstr Int [B], (1993) 53 (11) 5515.

ISSN: 0419-4217.

DOCUMENT TYPE:

(THESIS)

LANGUAGE:

English

FILE SEGMENT: ENTRY MONTH: Institute for Cell and Developmental Biology

199311

ENTRY DATE:

Entered STN: 19941107

Last Updated on STN: 19970509

Heparanase, an endo-beta-D-glucuronidase, has been associated with AB melanoma metastasis. Polyclonal antibodies directed against the murine N-terminal heparanase peptide detected a Mr of approx 97,000 protein upon SDS-polyacrylamide gel electrophoresis of mouse melanoma and human melanoma cell lysates. In an indirect immunocytochemical study, metastatic human A375-SM and mouse B16-BL6 melanoma cells were stained with the antiheparanase antibodies. Heparanase antigen was localized in the cytoplasm of permeabilized melanoma cells as well as at the cell surface of unpermeabilized cells. Immunohistochemical staining of frozen sections from syngeneic mouse organs containing micrometastases of B16-BL6 melanoma demonstrated heparanase localized in metastatic melanoma cells, but not in adjacent normal tissues. Similar studies using frozen sections of malignant melanomas resected from patients indicated that heparanase is localized in invading melanoma cells, but not in adjacent connective tissues. Monoclonal antibodies directed against murine heparanase were developed and characterized. Monoclonal antibody 10E5, an IgM, precipitated and inhibited the enzymatic activity of heparanase. A 2.6-kb cDNA was isolated from a human melanoma lambda gt11 cDNA library using the monoclonal antibody 10E5. Heparan sulfate cleavage activity was detected in the lysogen lysates from E coli Y1089 infected with the lambda qt11 cDNA and this activity was inhibited in the presence of 10-fold excess of heparin, a potent inhibitor of heparanase. The nucleotide sequence of the cDNA was determined and insignificant homology was found with the gene sequences currently known. The cDNA hybridized to a 3.2-3.4 kb mRNA in human A375 melanoma, WI-38 fibroblast, and THP-1 leukemia cells using Northern blots. Heparanase expression was examined using Western and Northern blots. In comparison to human A375-P melanoma cells, the quantity of 97,000 protein recognized by the polyclonal anti-heparanase antibodies doubled in the metastatic variant A375-SM cells and the quantity of 3.2-3.4 kb mRNA doubled in A375MetMix, a metastatic variant similar to A375-SM cells. In B16 murine melanoma cell, the intensity of the 97,000 protein increased more than 2 times comparing with B16-F1 cells. The extent in the increase of the protein and the $\ensuremath{\mathtt{mRNA}}$ levels is comparable to the change of heparanase activity observed in those cells. In summary, the studies suggest that (a) the N-terminus of the heparanase molecule in mouse and human is antigenically related; (b) heparanase antiqens are localized at the cell surface and in the cytoplasm of metastatic human and mouse melanoma cells; (c) heparanase antigens are localized in invasive and metastatic murine and human melanomas in vivo, but not in adjacent normal tissues; (d) heparanase molecule appeared to be differentially expressed at the transcriptional as well as at the translational level; and (e) the size of human heparanase mRNA is 3.2-3.4 kb. (Full text available from University Microfilms International, Ann Arbor, MI, as Order No. AAD93-07237)

=> d 17 ibib ab 1-2

L7 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER:

2002:51645 CAPLUS

DOCUMENT NUMBER:

136:97379

TITLE:

Protein and cDNA sequences of a second human heparanase, and splice variants thereof, with

a predominant expression in skeletal muscle, heart and

pancreas

INVENTOR(S):

David, Guido; Duerr, Joachim

PATENT ASSIGNEE(S):

Vlaams Interuniversitair Instituut voor Biotechnologie

Vzw, Belg.

SOURCE:

PCT Int. Appl., 54 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

LANGUAGE:

Patent English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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APPLICATION NO. DATE
           KIND DATE
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PATENT NO.
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_____ _____
                           WO 2001-EP8094 20010712
WO 2002004645 A2
                 20020117
                20021017
WO 2002004645
            Α3
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: EP 2000-202442 A 20000712

The present invention provides a novel human protein and its splicing variants, which has heparanase activity. The present invention relates to the field of carbohydrates and more specifically to the field of heparan sulfate proteoglycans. Several splice variants of said gene have been identified with a specific expression pattern in skeletal muscle, heart and pancreas.

ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2001:229058 CAPLUS

DOCUMENT NUMBER:

134:262849

TITLE:

Human heparanase-2, its

sequence, recombinant production, and use in identifying potential antagonists and/or agonists

Duecker, Klaus; Sirrenberg, Christian

INVENTOR(S): PATENT ASSIGNEE(S):

Merck Patent G.m.b.H., Germany PCT Int. Appl., 46 pp.

SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND DA'	TE	APPLICATION NO.	DATE
	- -			
WO 2001021814	A1 20	010329	WO 2000-EP8837	20000911
W: CA, J	•			
RW: AT, B	E, CH, CY, D	E, DK, ES, F	'I, FR, GB, GR, 1	E, IT, LU, MC, NL,
PT, S	<u> </u>			
EP 1214423	A1 20	020619	EP 2000-958531	20000911
R: AT, B	C, CH, DE, D	K, ES, FR, G	B, GR, IT, LI, I	U, NL, SE, MC, PT,
IE, F	CY CY			
JP 2003510053	T2 20	030318	JP 2001-525372	20000911
PRIORITY APPLN. IN	O.:	EP) 1999-118805 <i>I</i>	19990923
		EP	2000-114649	20000707

WO 2000-EP8837 W 20000911

AΒ The invention provides a cDNA mol. encoding a human protein believed to be heparanase-2, based on sequence homol. to known heparanases. The invention also provides polynucleotides that contain fragments of said cDNA mols. that can be used as hybridization probes or as primers for nucleic acid amplification. The invention further provides expression vectors comprising said cDNA mols., host cells transformed with said vectors for the recombinant prodn. of human heparanase-2. Still further, the invention provides for

the use of said human heparanase-2

polypeptides in identifying compds. that may be antagonists and/or agonists, which may be potentially useful in therapy. Finally, the invention provides a fusion protein consisting of said heparanase -2 fused to a Ig Fc region, and antibodies specific for heparanase-2. The cDNA sequence, as well as the corresponding amino acid sequence of human heparanase-2 are claimed. The invention used reverse transcription-polymerase chain reaction (RT-PCR) to show the expression of heparanase-2 gene in various tissues and tumors, and showed the expression of heparanase-2 in transformed 293 human kidney fibroblasts.

REFERENCE COUNT:

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

WEST

Generate Collection

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Search Results - Record(s) 1 through 4 of 4 returned.

L2: Entry 1 of 4

File: PGPB

May 1, 2003

PGPUB-DOCUMENT-NUMBER: 20030083254

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030083254 A1

TITLE: Substances

PUBLICATION-DATE: May 1, 2003

INVENTOR - INFORMATION:

NAME CITY STATE COUNTRY RULE-47

McKenzie, Edward Alexander Abingdon GB Stamps, Alasdair Craig Abingdon GB Terrett, Jonathan Alexander Abingdon GB Tyson, Kerry Louise Abingdon GB

US-CL-CURRENT: 514/12; 530/324

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw Desc Image

☐ 2. Document ID: US 20020064853 A1

L2: Entry 2 of 4

File: PGPB

May 30, 2002

PGPUB-DOCUMENT-NUMBER: 20020064853

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020064853 A1

TITLE: Heparanase II, a novel human heparanase paralog

PUBLICATION-DATE: May 30, 2002

INVENTOR - INFORMATION:

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US-CL-CURRENT: 435/200; 435/18, 435/325, 435/69.1, 536/23.2

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw Desc Image

L2: Entry 3 of 4

File: USPT

May 14, 2002

US-PAT-NO: 6387643

DOCUMENT-IDENTIFIER: US 6387643 B1

** See image for Certificate of Correction **

TITLE: Human platelet heparanase polypeptides, polynucleotide molecules that encode them, and methods for the identification of compounds that alter heparanase activity

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	human heparanase	50	<u>L1</u>

END OF SEARCH HISTORY